IN THE CLAIMS

COMPLETE LISTING OF ALL CLAIMS, WITH MARKINGS AND STATUS IDENTIFIERS

In the revised claim set appearing below, currently amended claims have deletions shown by strikethrough or brackets [[]], and additions shown by underlining. This listing of claims will replace all prior versions and listings of the claims in the application.

Listing of Claims:

- 1. (canceled)
- 2. (currently amended) The method A peptide according to claim [[1]] 6 or 9, wherein said peptide of formula (I) is a peptide wherein

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A¹ is L-Phe, D-Phe, L-Cpa or D-Cpa;
A³ is L-Tyr, L-Trp or L-3-Pal;
A⁴ is D-Trp;
A⁶ is β-Ala or Gaba;
Aⁿ is L-Cys;
Aⁿ is L-Thr, L-Trp, L-Leu or L-Nal; and
R² and R³ are each H;
or a pharmaceutically acceptable salt thereof.
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3. (currently amended) The method A peptide according to claim [[2]] 6 or 9, wherein said peptide is of the formula

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Cpa-cyclo(D-Cys-3-Pal-D-Trp-Lys-Gaba-Cys)-Nal-NH<sub>2</sub>;
Cpa-cyclo(D-Cys-3-Pal-D-Trp-Lys-β-Ala-Cys)-Nal-NH<sub>2</sub>;
Phe-cyclo(D-Cys-3-Pal-D-Trp-Lys-Gaba-Cys)-Nal-NH<sub>2</sub>;
Phe-cyclo(D-Cys-Tyr-D-Trp-Lys-Gaba-Cys)-Nal-NH<sub>2</sub>;
Phe-cyclo(D-Cys-Trp-D-Trp-Lys-Gaba-Cys)-Nal-NH<sub>2</sub>;
Phe-cyclo(D-Cys-Tyr-D-Trp-Lys-Gaba-Cys)-Trp-NH<sub>2</sub>;
D-Phe-cyclo(D-Cys-Tyr-D-Trp-Lys-Gaba-Cys)-Nal-NH<sub>2</sub>;
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D-Phe-cyclo(D-Cys-Tyr-D-Trp-Lys-Gaba-Cys)-Leu-NH₂; or Phe-cyclo-(D-Cys-Tyr-D-Trp-Lys-Gaba-Cys)-Thr-NH₂; or a pharmaceutically acceptable salt thereof.

4. (currently amended) The method A peptide according to claim [[3]] 6 or 9, wherein said peptide is of the formula

Cpa-cyclo(D-Cys-3-Pal-D-Trp-Lys-Gaba-Cys)-Nal-NH₂; or Cpa-cyclo(D-Cys-3-Pal-D-Trp-Lys-β-Ala-Cys)-Nal-NH₂; or a pharmaceutically acceptable salt thereof.

- 5. (currently amended) [[A]] The method according to claim 6 or 9, wherein said peptide or pharmaceutically acceptable salt thereof is in the form of a pharmaceutical composition useful for eliciting a somatostatin agonist response in a human or other animal which comprises an effective amount of a peptide of formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.
- 6. (previously presented) A method of eliciting a somatostatin agonist response in a human or other animal in need thereof, which comprises administering an effective amount of a peptide of formula (I) X-A¹-cyclo(D-Cys-A³-A⁴-Lys-A⁶-A³)-A³-Y,

(I)

or a pharmaceutically acceptable salt thereof, wherein

A¹ and A³ are each independently the D- or L-isomer of an amino acid selected from the group consisting of Phe, Tyr, Tyr(I), Trp, 3-Pal, 4-Pal, Cpa and Nal;

A⁴ is L-Trp, D-Trp, L-\u00a3-methyl-Trp or D-\u00a3-methyl-Trp;

 A^6 is -NH-(CHR¹)_n-CO-, where n is 2, 3, or 4;

 A^7 is L- or D-Cys;

A⁸ is the D- or L-isomer of an amino acid selected from the group consisting of Phe, Tyr, Tyr(I), Trp, Nal, Cpa, Val, Leu, Ile, Ser and Thr;

Y is NR²R³ where R² and R³ are each independently H or (C₁-C₅)alkyl;

 R^1 is selected from the group consisting H, (C_1-C_4) alkyl and $-CH_2$ -aryl; wherein said aryl is an optionally substituted moiety selected from the group consisting of phenyl, 1-naphthyl, and 2-naphthyl, wherein said optionally substituted moiety is optionally substituted with one or more substituents each independently selected from the group consisting of (C_{1-6}) alkyl, (C_{2-6}) alkenyl, (C_{2-6}) alkynyl, aryl, aryl (C_{1-6}) alkyl, (C_{1-6}) alkoxy, $-N(R^4R^5)$, -COOH, $-CON(R^4R^5)$, halo, -OH, -CN, and $-NO_2$;

 R^4 and R^5 each is, independently for each occurrence, H or (C_{1-3}) alkyl; where the Cys of A^2 is bonded to the Cys of A^7 by a di-sulfide bond formed from the thiol groups of each Cys,

to the human or other animal.

- 7. (canceled)
- 8. (canceled)
- 9. (currently amended) A method of inhibiting the secretion of growth hormone, insulin, glucagon or pancreatic exocrine secretion in a human or other animal in need thereof, which comprises administering a peptide of formula (I)

$$X-A^1$$
-cyclo(D-Cys- A^3 - A^4 -Lys- A^6 - A^7)- A^8 - Y ,
(I)

or a pharmaceutically acceptable salt thereof, wherein

$$X \text{ is } H,$$
 HO(CH₂)₂-N N-(CH₂)-CO- or HO(CH₂)₂-N N-(CH₂)₂-SO₂-

A¹ and A³ are each independently the D- or L-isomer of an amino acid selected from the group consisting of Phe, Tyr, Tyr(I), Trp, 3-Pal, 4-Pal, Cpa and Nal;

A⁴ is L-Trp, D-Trp, L-\beta-methyl-Trp or D-\beta-methyl-Trp;

 A^6 is -NH-(CHR¹)_n-CO-, where n is 2, 3, or 4;

A⁷ is L- or D-Cys;

A⁸ is the D- or L-isomer of an amino acid selected from the group consisting of Phe, Tyr, Tyr(I), Trp, Nal, Cpa, Val, Leu, Ile, Ser and Thr;

Y is NR²R³ where R² and R³ are each independently H or (C₁-C₅)alkyl;

 R^1 is selected from the group consisting H, (C_1-C_4) alkyl and -CH₂-aryl; wherein said aryl is an optionally substituted moiety selected from the group consisting of phenyl, 1-naphthyl, and 2-naphthyl, wherein said optionally substituted moiety is optionally substituted with one or more substituents each independently selected from the group consisting of (C_{1-6}) alkyl, (C_{2-6}) alkenyl, (C_{2-6}) alkynyl, aryl, aryl (C_{1-6}) alkyl, (C_{1-6}) alkoxy, -N((R^4R^5)), -COOH, -CON((R^4R^5)), halo, -OH, -CN, and -NO₂;

 R^4 and R^5 each is, independently for each occurrence, H or (C_{1-3}) alkyl; where the Cys of A^2 is bonded to the Cys of A^7 by a di-sulfide bond formed from the thiol groups of each Cys,

to said human or other animal.

- 10. (canceled)
- 11. (canceled)